

Rec'd PCT/PTO 20 DEC 2004  
PCT/EP 03/06097

04.07.03



Europäisches  
Patentamt

European  
Patent Office

Office européen  
des brevets

REC'D 04 AUG 2003

WIPO

PCT

Bescheinigung

Certificate

Attestation

Die angehefteten Unterla-  
gen stimmen mit der  
ursprünglich eingereichten  
Fassung der auf dem näch-  
sten Blatt bezeichneten  
europäischen Patentanmel-  
dung überein.

The attached documents  
are exact copies of the  
European patent application  
described on the following  
page, as originally filed.

Les documents fixés à  
cette attestation sont  
conformes à la version  
initialement déposée de  
la demande de brevet  
européen spécifiée à la  
page suivante.

Patentanmeldung Nr. Patent application No. Demande de brevet n°

02014214.7

**PRIORITY  
DOCUMENT**  
SUBMITTED OR TRANSMITTED IN  
COMPLIANCE WITH RULE 17.1(a) OR (b)

Der Präsident des Europäischen Patentamts;  
Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets  
p.o.

R C van Dijk



Europäisches  
Patentamt

YCT/EP 03/06097  
European  
Patent Office

Office européen  
des brevets

04.03.03

Anmeldung Nr:  
Application no.: 02014214.7  
Demande no:

Anmeldetag:  
Date of filing: 26.06.02  
Date de dépôt:

Anmelder/Applicant(s)/Demandeur(s):

Basell Poliolefine Italia S.p.A.  
Via Pergolesi, 25  
20124 Milano  
ITALIE

Bezeichnung der Erfindung/Title of the invention/Titre de l'invention:  
(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.  
If no title is shown please refer to the description.  
Si aucun titre n'est indiqué se référer à la description.)

Impact-resistant polyolefin compositions

In Anspruch genommene Priorität(en) / Priority(ies) claimed /Priorité(s)  
revendiquée(s)  
Staat/Tag/Aktenzeichen/State/Date/File no./Pays/Date/Numéro de dépôt:

Internationale Patentklassifikation/International Patent Classification/  
Classification internationale des brevets:

C08L23/00

Am Anmeldetag benannte Vertragstaaten/Contracting states designated at date of  
filing/Etats contractants désignées lors du dépôt:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR

EPO - Munich  
68  
26. Juni 2002**IMPACT-RESISTANT POLYOLEFIN COMPOSITIONS**

The present invention concerns polyolefin compositions comprising a crystalline propylene polymer component selected from propylene-ethylene and/or other  $\alpha$ -olefin random copolymers, and a copolymer of ethylene with  $C_4$ - $C_{10}$   $\alpha$ -olefins.

The compositions of the present invention can be easily converted into various kinds of finished or semi-finished articles, in particular by using injection-molding techniques, as they possess relatively high values of melt flow rate (MFR). In addition, as they show substantially no stress whitening when bending a plaque 1 mm thick, the said compositions can be used for several applications, including toys and housewares, in particular for those articles that need with impact resistance at low temperatures without failure of the articles. The said articles can advantageously be used in the food-contact applications, examples of which are food containers suitable for freezers.

Compositions comprising polypropylene and a rubbery phase formed by an elastomeric copolymer of ethylene with  $\alpha$ -olefins are already known in the art, and described in particular in European patents 170 255 and 373 660, and in WO 01/19915. Said compositions present impact resistance and, in the case of European patent 373 660 and WO 01/19915, transparency values interesting for many applications, however the overall balance of properties is still not totally satisfactory in the whole range of possible applications, in view of the high standards required by the market. Therefore there is a continuous demand for compositions of this kind with improved properties.

A new and valuable balance of properties has now been achieved by the polyolefin compositions of the present invention, comprising (percent by weight):

- 1) 55-80% of a crystalline propylene homopolymer or copolymer containing up to 15% of ethylene and/or  $C_4$ - $C_{10}$   $\alpha$ -olefin(s) and having an MFR value of at least 15 g/10 min; and
- 2) 20-45% of a copolymer of ethylene with one or more  $C_4$ - $C_{10}$   $\alpha$ -olefin(s) containing from 10 to 40% of said  $C_4$ - $C_{10}$   $\alpha$ -olefin(s);

said compositions having values of MFR (230 °C, 2.16 Kg) equal to or higher than 15 g/10 min, a total content of ethylene of 20% or more, a total content of  $C_4$ - $C_{10}$   $\alpha$ -olefin(s) of 4.5% or more, a ratio of the total content of ethylene to the total content of  $C_4$ - $C_{10}$   $\alpha$ -olefin(s) of 2.3 or more, and an intrinsic viscosity value of the fraction soluble in xylene at room temperature of 1.7 dl/g or less, preferably of 1.5 dl/g or less.

From the above definitions it is evident that the term "copolymer" includes polymers containing more than one kind of comonomers.

The compositions of the present invention provide in particular a combination of very high flowability and high impact resistance (in terms of ductile/brittle transition temperature and Izod impact resistance) and high transparency.

The preferred polyolefin compositions are flexible polyolefin compositions comprising (percent by weight):

- 1) 55-75%, preferably 55-70%, of a crystalline propylene homopolymer or copolymer containing up to 15% of ethylene and/or C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) and having an MFR value of from 15 to 80 g/10 min; and
- 2) 25-45%, preferably 30-45%, of a copolymer of ethylene with one or more C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) containing from 15 to 40% of said C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s);

said compositions having values of MFR (230 °C, 2.16 Kg) equal to or higher than 15 g/10 min, a total content of ethylene of 20% or more, a total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 6% or more, a ratio of the total content of ethylene to the total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 2.3 or more, a total fraction soluble in xylene at room temperature of 18 wt% or higher, preferably at least 20 wt%, and an intrinsic viscosity value of the fraction soluble in xylene at room temperature of 1.7 dl/g or less, preferably of 1.5 dl/g or less.

The compositions of the present invention have preferably a MFR value in the range of from 15 g/10 to 40 g/10 min.

Particularly preferred features for the compositions of the present invention are:

- content of polymer insoluble in xylene at room temperature (23 °C) (substantially equivalent to the Isotacticity Index) for component 1): not less than 90%, in particular not less than 93%, said percentages being by weight and referred to the weight of component 1);
- a total content of ethylene from 20% to 40% by weight;
- a total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) from 6% to 15% by weight;
- a Flexural Modulus from less than 770 MPa, preferably higher than 600 MPa, more preferably higher than 650 MPa;
- fraction soluble in xylene at room temperature: less than 35%, more preferably less than 30% by weight;
- intrinsic viscosity of the fraction soluble in xylene at room temperature in the range

from 0.8 to 1.5 dl/g.

The ductile/brittle transition temperature is generally equal to or lower than  $-35^{\circ}\text{C}$ , the lower limit being indicatively of about  $-60^{\circ}\text{C}$ .

The said  $\text{C}_4\text{-C}_{10}$   $\alpha$ -olefins, that are or may be present as comonomers in the components and fractions of the compositions of the present invention, are represented by the formula  $\text{CH}_2=\text{CHR}$ , wherein R is an alkyl radical, linear or branched, with 2-8 carbon atoms or an aryl (in particular phenyl) radical.

Examples of said  $\text{C}_4\text{-C}_{10}$   $\alpha$ -olefins are 1-butene, 1-pentene, 1-hexene, 4-methyl-1-pentene and 1-octene. Particularly preferred is 1-butene.

The compositions of the present invention can be prepared by a sequential polymerization, comprising at least two sequential steps, wherein components 1) and 2) are prepared in separate subsequent steps, operating in each step, except the first step, in the presence of the polymer formed and the catalyst used in the preceding step. The catalyst is added only in the first step, however its activity is such that it is still active for all the subsequent steps.

Preferably component 1) is prepared before component 2).

The polymerization, which can be continuous or batch, is carried out following known techniques and operating in liquid phase, in the presence or not of inert diluent, or in gas phase, or by mixed liquid-gas techniques. Preferably both components 1) and 2) are prepared in gas phase.

Reaction time, pressure and temperature relative to the two steps are not critical, however it is best if the temperature is from 20 to  $100^{\circ}\text{C}$ . The pressure can be atmospheric or higher.

The regulation of the molecular weight is carried out by using known regulators, hydrogen in particular.

Such polymerization is preferably carried out in the presence of stereospecific Ziegler-Natta catalysts. An essential component of said catalysts is a solid catalyst component comprising a titanium compound having at least one titanium-halogen bond, and an electron-donor compound, both supported on a magnesium halide in active form. Another essential component (co-catalyst) is an organoaluminum compound, such as an aluminum alkyl compound.

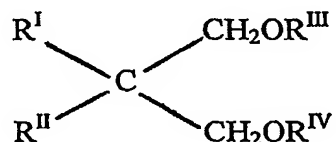
An external donor is optionally added.

The catalysts generally used in the process of the invention are capable of producing polypropylene with an isotactic index greater than 90%, preferably greater than 95%. Catalysts having the above mentioned characteristics are well known in the patent literature; particularly advantageous are the catalysts described in US patent 4,399,054 and European patent 45977.

The solid catalyst components used in said catalysts comprise, as electron-donors (internal donors), compounds selected from the group consisting of ethers, ketones, lactones, compounds containing N, P and/or S atoms, and esters of mono- and dicarboxylic acids.

Particularly suitable electron-donor compounds are phthalic acid esters, such as diisobutyl, dioctyl, diphenyl and benzylbutyl phthalate.

Other electron-donors particularly suitable are 1,3-diethers of formula:



wherein R<sup>I</sup> and R<sup>II</sup> are the same or different and are C<sub>1</sub>-C<sub>18</sub> alkyl, C<sub>3</sub>-C<sub>18</sub> cycloalkyl or C<sub>7</sub>-C<sub>18</sub> aryl radicals; R<sup>III</sup> and R<sup>IV</sup> are the same or different and are C<sub>1</sub>-C<sub>4</sub> alkyl radicals; or are the 1,3-diethers in which the carbon atom in position 2 belongs to a cyclic or polycyclic structure made up of 5, 6 or 7 carbon atoms and containing two or three unsaturations.

Ethers of this type are described in published European patent applications 361493 and 728769.

Representative examples of said diethers are 2-methyl-2-isopropyl-1,3-dimethoxypropane, 2,2-diisobutyl-1,3-dimethoxypropane, 2-isopropyl-2-cyclopentyl-1,3-dimethoxypropane, 2-isopropyl-2-isoamyl-1,3-dimethoxypropane, 9,9-bis (methoxymethyl) fluorene.

The preparation of the above mentioned catalyst components is carried out according to various methods.

For example, a MgCl<sub>2</sub>·nROH adduct (in particular in the form of spheroidal particles) wherein n is generally from 1 to 3 and ROH is ethanol, butanol or isobutanol, is reacted with an excess of TiCl<sub>4</sub> containing the electron-donor compound. The reaction temperature is generally from 80 to 120 °C. The solid is then isolated and reacted once more with TiCl<sub>4</sub>, in the presence or absence of the electron-donor compound, after which it is separated and washed with aliquots of a hydrocarbon until all chlorine ions have disappeared.

In the solid catalyst component the titanium compound, expressed as Ti, is generally present in an amount from 0.5 to 10% by weight. The quantity of electron-donor compound which remains fixed on the solid catalyst component generally is 5 to 20% by moles with respect to the magnesium dihalide.

The titanium compounds which can be used for the preparation of the solid catalyst component are the halides and the halogen alcoholates of titanium. Titanium tetrachloride is the preferred compound.

The reactions described above result in the formation of a magnesium halide in active form. Other reactions are known in the literature, which cause the formation of magnesium halide in active form starting from magnesium compounds other than halides, such as magnesium carboxylates.

The Al-alkyl compounds used as co-catalysts comprise the Al-trialkyls, such as Al-triethyl, Al-triisobutyl, Al-tri-n-butyl, and linear or cyclic Al-alkyl compounds containing two or more Al atoms bonded to each other by way of O or N atoms, or  $\text{SO}_4$  or  $\text{SO}_3$  groups. The Al-alkyl compound is generally used in such a quantity that the Al/Ti ratio be from 1 to 1000.

The electron-donor compounds that can be used as external donors include aromatic acid esters such as alkyl benzoates, and in particular silicon compounds containing at least one Si-OR bond, where R is a hydrocarbon radical. Examples of silicon compounds are  $(\text{tert-butyl})_2\text{Si}(\text{OCH}_3)_2$ ,  $(\text{cyclohexyl})(\text{methyl})\text{Si}(\text{OCH}_3)_2$ ,  $(\text{phenyl})_2\text{Si}(\text{OCH}_3)_2$  and  $(\text{cyclopentyl})_2\text{Si}(\text{OCH}_3)_2$ . 1,3-diethers having the formulae described above can also be used advantageously. If the internal donor is one of these diethers, the external donors can be omitted.

The catalysts can be pre-contacted with small amounts of olefins (prepolymerization).

Other catalysts that may be used in the process according to the present invention are metallocene-type catalysts, as described in USP 5,324,800 and EP-A-0 129 368; particularly advantageous are bridged bis-indenyl metallocenes, for instance as described in USP 5,145,819 and EP-A-0 485 823. Another class of suitable catalysts are the so-called constrained geometry catalysts, as described in EP-A-0 416 815 (Dow), EP-A-0 420 436 (Exxon), EP-A-0 671 404, EP-A-0 643 066 and WO 91/04257. These metallocene compounds may be used in particular to produce the copolymers (a) and (b).

The compositions of the present invention can also be obtained by preparing separately

the said components 1) and 2), by operating with the same catalysts and substantially under the same polymerization conditions as previously explained (except that a wholly sequential polymerization process will not be carried out, but the said components will be prepared in separate polymerization steps) and then mechanically blending said components in the molten or softened state. Conventional mixing apparatuses, like screw extruders, in particular twin screw extruders, can be used.

The compositions of the present invention can also contain additives commonly employed in the art, such as antioxidants, light stabilizers, heat stabilizers, nucleating agents, colorants and fillers.

In particular, the addition of nucleating agents brings about a considerable improvement in important physical-mechanical properties, such as flexural modulus, Heat Distortion Temperature (HDT), tensile strength at yield and transparency.

Typical examples of nucleating agents are the p-tert.-butyl benzoate and the 1,3- and 2,4-dibenzylidenesorbitols.

The nucleating agents are preferably added to the compositions of the present invention in quantities ranging from 0.05 to 2% by weight, more preferably from 0.1 to 1% by weight with respect to the total weight.

The addition of inorganic fillers, such as talc, calcium carbonate and mineral fibers, also brings about an improvement to some mechanical properties, such as flexural modulus and HDT. Talc can also have a nucleating effect.

The particulars are given in the following examples, which are given to illustrate, without limiting, the present invention.

### Examples 1-3

In the following examples polyolefin compositions according to the present invention are prepared by sequential polymerization.

The solid catalyst component used in polymerization is a highly stereospecific Ziegler-Natta catalyst component supported on magnesium chloride, containing about 2.5% by weight of titanium and diisobutylphthalate as internal donor, prepared by analogy with the method described in Example 1 of European published patent application 674991.

### CATALYST SYSTEM AND PREPOLYMERIZATION TREATMENT

Before introducing it into the polymerization reactors, the solid catalyst component described above is contacted at 100°C for 30 minutes with aluminum triethyl (TEAL) and



dicyclopentylmethoxysilane (DCPMS), in a TEAL/DCPMS weight ratio equal to about 4 and in such quantity that the TEAL/Ti molar ratio be equal to 65.

The catalyst system is then subjected to prepolymerization by maintaining it in suspension in liquid propylene at 20 °C for about 20 minutes before introducing it into the first polymerization reactor.

### POLYMERIZATION

The polymerization is carried out in continuous in a series of two gas phase reactors equipped with devices for the transfer of the product coming from the reactor immediately preceding to the one immediately following.

In gas phase the hydrogen and the monomer(s) are analyzed in continuous and fed in such a manner that the desired concentration be maintained constant.

Into a first gas phase polymerization reactor a propylene/ethylene copolymer is produced by feeding in a continuous and constant flow the prepolymerized catalyst system, hydrogen (used as molecular weight regulator) and propylene and ethylene monomers in the gas state, thus obtaining component 1).

The polymer produced in the first reactor is discharged in the second reactor where an ethylene/butene copolymer is produced by feeding the monomer(s) and hydrogen in proper molar ratios, thus obtaining component 2).

Then the polymer particles are introduced in a rotating drum, where they are mixed with 0.05% by weight of paraffinic oil, 0.05% by weight of sodium stearate, 0.15% by weight of Irganox® B215 (1 weight part of pentaerythryl-tetrakis[3(3,5-di-tert-butyl-4-hydroxyphenyl)] mixed with 1 weight part of tris(2,4-ditert-butylphenyl) phosphite) and 0.2% by weight of Millad® 3988 3,4-dimethylbenzylidene sorbitol.

Then the polymer particles are introduced in a twin screw extruder Berstorff™ ZE 25 (length/diameter ratio of screws: 33) and extruded under nitrogen atmosphere in the following conditions:

Rotation speed:	250 rpm;
Extruder output:	6-20 kg/hour;
Melt temperature:	200-250 °C.

The data relating to the final polymer compositions reported in table 1 and 2 are obtained from measurements carried out on the so extruded polymers.

The data shown in the tables are obtained by using the following test methods.

- **Molar ratios of the feed gases**

Determined by gas-chromatography..

- **Ethylene and 1-butene content of the polymers**

Determined by I.R. spectroscopy

- **Melt Flow Rate (MFR)**

Determined according to ASTM D 1238, condition L (MFR“L”).

- **Xylene soluble and insoluble fractions**

Determined as follows.

2.5 g of polymer and 250 ml of xylene are introduced in a glass flask equipped with a refrigerator and a magnetical stirrer. The temperature is raised in 30 minutes up to the boiling point of the solvent. The so obtained clear solution is then kept under reflux and stirring for further 30 minutes. The closed flask is then kept for 30 minutes in a bath of ice and water and in thermostatic water bath at 25 °C for 30 minutes as well. The so formed solid is filtered on quick filtering paper. 100 ml of the filtered liquid is poured in a previously weighed aluminum container which is heated on a heating plate under nitrogen flow, to remove the solvent by evaporation. The container is then kept in an oven at 80 °C under vacuum until constant weight is obtained. The weight percentage of polymer soluble in xylene at room temperature is then calculated. The percent by weight of polymer insoluble in xylene at room temperature is considered the Isotacticity Index of the polymer. This value corresponds substantially to the Isotacticity Index determined by extraction with boiling n-heptane, which by definition constitutes the Isotacticity Index of polypropylene.

- **Intrinsic Viscosity (I.V.)**

Determined in tetrahydronaphthalene at 135 °C.

- **Flexural Modulus**

Determined according to ISO 178.

- **Ductile/Brittle transition temperature (D/B)**

Determined according to internal method MA 17324, available upon request.

According to this method, the bi-axial impact resistance is determined through impact with an automatic, computerised striking hammer.

The circular test specimens are obtained by cutting with circular hand punch (32 mm

diameter). They are conditioned for at least 12 hours at 23°C and 50 RH and then placed in a thermostatic bath at testing temperature for 1 hour.

The force-time curve is detected during impact of a striking hammer (5.3 kg, hemispheric punch with a ½" diameter) on a circular specimen resting on a ring support. The machine used is a CEAST 6758/000 type model no. 2.

D/B transition temperature means the temperature at which 50% of the samples undergoes fragile break when submitted to the said impact test.

#### **Preparation of the plaque specimens**

Plaques for D/B measurement, having dimensions of 127x127x1.5 mm are prepared according to internal method MA 17283; plaques for haze measurement, 1 mm thick, are prepared by injection moulding according to internal method MA 17335 with injection time of 1 second, temperature of 230 °C, mould temperature of 40 °C, description of all the said methods being available upon request.

#### **Method MA 17283**

The injection press is a Negri Bossi™ type (NB 90) with a clamping force of 90 tons. The mould is a rectangular plaque (127x127x1.5mm).

The main process parameters are reported below:

Back pressure (bar):	20
Injection time (s):	3
Maximum Injection pressure (MPa):	14
Hydraulic injection pressure (MPa):	6-3
First holding hydraulic pressure (MPa):	4±2
First holding time (s):	3
Second holding hydraulic pressure (MPa):	3±2
Second holding time (s):	7
Cooling time (s):	20
Mould temperature (°C):	60

The melt temperature is between 220 and 280 °C.

#### **Method MA 17335**

The injection press is a Battenfeld™ type BA 500CD with a clamping force of 50 tons. The insert mould leads to the moulding of two plaques (55x60x1 or 1.5 mm each).

- **Haze on plaque**

Determined according to internal method MA 17270, available upon request.

The plaques are conditioned for 12 to 48 hours at R.H.  $50\pm 5\%$  and  $23\pm 1^\circ\text{C}$ .

The apparatus used is a Hunter™ D25P-9 colorimeter. The measurement and computation principle are given in the norm ASTM-D1003.

The apparatus is calibrated without specimen, the calibration is checked with a haze standard. The haze measurement is carried out on five plaques.

- **Izod impact strength (notched)**

Determined according to ISO180/1A.

**Comparative example 1c**

Example 1 is repeated except that the polymerisation is carried out in a series of three reactors. Into the first reactor a crystalline propylene-ethylene copolymer is produced feeding the monomers and hydrogen in proper molar ratios (component (A')). The copolymer thus produced is discharged into the second reactor where a propylene-ethylene copolymer is produced by feeding the monomers and hydrogen in proper molar ratios (component (A'')).

The copolymer produced in the second reactor is discharged in a continuous flow and, after having being purged of unreacted monomers, is introduced in a continuous flow into the third gas phase reactor, together with quantitatively constant flows of hydrogen and ethylene and 1-butene monomers in the gas state. Component (B) is thus obtained.

Polymerisation conditions, molar ratios, composition and properties of the copolymers obtained are shown in table 2. The comparative composition shows a value of flexural modulus in the same range as the one of the compositions of the present invention, value which is obtained only thanks to a crystalline polymer moiety of the matrix having a low flowability.

In comparison with the comparative composition, the compositions according to the present invention have a comparable or even better stiffness and better impact resistance in terms of ductile/brittle transition temperature in spite of remarkably higher MFR values that improve workability as it generally affects stiffness and impact resistance.

TABLE 1

Example and comparative example		1	2	3	1c
1 <sup>st</sup> Gas Phase Reactor - crystalline propylene-ethylene copolymer					
Temperature	°C	80	80	80	80
Pressure	MPa	-	-	-	1.8
H <sub>2</sub> /C <sub>3</sub>	mol	-	-	-	0.002
C <sub>2</sub> /(C <sub>2</sub> +C <sub>3</sub> )	mol	-	-	-	0.019
MFR "L"	g/10'	41	52	24.4	1.2
Ethylene content in the copolymer	wt%	2.1	2.0	2.4	2.6
Xylene soluble fraction	wt%	3.6	-	4.1	-
Split	wt%	66	68	69	39
2 <sup>nd</sup> Gas Phase Reactor - crystalline propylene-ethylene copolymer					
Split	wt%	0	0	0	39
MFR "L" (total)	g/10'	-	-	-	11.7
Ethylene content in the copolymer	wt%	-	-	-	2.6
H <sub>2</sub> /C <sub>3</sub>	mol	-	-	-	0.419
Xylene-soluble fraction (total)	wt%	-	-	-	96.5
2 <sup>nd</sup> /3 <sup>rd</sup> Gas Phase Reactor - ethylene-butene-1 copolymer rubber					
Temperature	°C	75	75	70	70
Pressure	MPa	-	-	-	1.6
H <sub>2</sub> /C <sub>2</sub>	mol			-	0.466
C <sub>4</sub> /(C <sub>4</sub> +C <sub>2</sub> )	mol	0.55	0.55	0.51	0.35
Split	wt%	34	32	31	22
Butene-1 in the rubber	wt%	27	24	25.8	23.6
Xylene soluble fraction	wt%	65	60	64	-

TABLE 2

Example and comparative example		1	2	3	1
MFR "L"	g/10'	32.5	28.2	19.4	9.4
Xylene-soluble fraction	wt%	24.6	-	22.6	13.6
I.V. of xylene-soluble fraction	dl/g	1.05	-	1.09	1.29
Ethylene content	wt%	26.0	25.7	25.4	18.4
Butene-1 content	wt%	9.1	7.7	8.0	5.2
Flexural modulus	MPa	671	757	760	1015
D/B transition temperature	°C	-53	-49	-50	-22
Izod impact resistance at 23° C	kJ/m <sup>2</sup>	-	-	37.9	190 <sup>1)</sup>
Haze, 1 mm plaque	%	35	39.5	23.7	13.3

<sup>1)</sup> Expressed in J/m; 190 J/m corresponds to about 15.1 kJ/m<sup>2</sup>.

Notes to the tables.

Split = weight fraction of polymer produced in the specified reactor;  $C_2^-$  = ethylene;  $C_4^-$  = butene;  $H_2/C_2^-$  = molar ratio of fed hydrogen to fed ethylene;  $C_2^-/(C_2^-+C_3^-)$  = molar ratio of fed ethylene to fed ethylene plus fed propylene;  $C_4^-/(C_4^-+C_2^-)$  = molar ratio of fed butene to fed butene plus fed ethylene.

## CLAIMS

Munich  
68  
26. Juni 2002

1. Polyolefin compositions comprising (percent by weight):
  - 1) 55-80% of a crystalline propylene homopolymer or copolymer containing up to 15% of ethylene and/or C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) and having an MFR value (230 °C, 2.16 Kg) of at least 15 g/10 min; and
  - 2) 20-45% of a copolymer of ethylene with one or more C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) containing from 10 to 40% of said C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s);said compositions having values of MFR (230 °C, 2.16 Kg) equal to or higher than 15 g/10 min, a total content of ethylene of 20% or more, a total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 4.5% or more, a ratio of the total content of ethylene to the total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 2.3 or more, and an intrinsic viscosity value of the fraction soluble in xylene at room temperature of 1.7 dl/g or less.
2. The polyolefin compositions according to claim 1 comprising (percent by weight):
  - 1) 55-75%, preferably 55-70%, of a crystalline propylene homopolymer or copolymer containing up to 15% of ethylene and/or C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) and having an MFR value of from 15 to 80 g/10 min; and
  - 2) 25-45%, preferably 30-45%, of a copolymer of ethylene with one or more C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) containing from 20 to 40% of said C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s);said compositions having values of MFR (230 °C, 2.16 Kg) equal to or higher than 15 g/10 min, a total content of ethylene of 20% or more, a total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 6% or more, a ratio of the total content of ethylene to the total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 2.3 or more, a total fraction soluble in xylene at room temperature of 18 wt% or higher, and an intrinsic viscosity value of the fraction soluble in xylene at room temperature of 1.7 dl/g or less.
3. The polyolefin compositions of claim 1, having MFR values equal to or higher than 30 g/10 min.
4. The polyolefin compositions of claim 1, wherein the intrinsic viscosity of the fraction soluble in xylene at room temperature is in the range from 0.8 to 1.5 dl/g.
5. The polyolefin compositions of claim 1, wherein the content of polymer soluble in xylene at room temperature is higher than 20%.
6. The polyolefin compositions of claim 1, having a ductile/brittle transition temperature equal to or lower than -35 °C.

7. A process for producing the polyolefin compositions of claim 1, carried out in at least two sequential steps, wherein in at least one polymerization step the relevant monomer(s) are polymerized to form component 1) and in the other step the relevant monomers are polymerized to form component 2), operating in each step, except the first step, in the presence of the polymer formed and the catalyst used in the preceding step.
8. The process of claim 6, wherein the polymerization catalyst is a stereospecific Ziegler-Natta catalyst comprising, as catalyst-forming components, a solid component comprising a titanium compound having at least one titanium-halogen bond and an electron-donor compound, both supported on a magnesium halide in active form, and an organoaluminum compound.
9. The process of claim 6, wherein both components 1) and 2) are prepared in gas phase.
10. Injection moulded articles comprising the polyolefin compositions of claim 1.



EPO · Munich  
68  
26. Juni 2002

# ABSTRACT

Polyolefin compositions comprise 1) 55-80 wt% of a crystalline propylene homopolymer or copolymer containing up to 15 wt% of ethylene and/or C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) and having an MFR value of at least 15 g/10 min and 2) 20-45 wt% of a copolymer of ethylene with one or more C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) containing from 10 to 4 wt% of said C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s). The said compositions have values of MFR equal to or higher than 15 g/10 min, a total content of ethylene of 20% or more, a total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 4.5% or more, a ratio of the total content of ethylene to the total content of C<sub>4</sub>-C<sub>10</sub>  $\alpha$ -olefin(s) of 2.3 or more, and an intrinsic viscosity value of the fraction soluble in xylene at room temperature of 1.7 dl/g or less.